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EEE TOANS	EE TRANSMITTAL		Complete if Known			
FEE IRANS	DIVITIAL	Application Number	09/699,723			
for EV 1	<b>1</b> 000	Filing Date	October 30, 2000			
for FY 2004  Effective 10/01/2003. Patent fees are subject to annual revision.		First Named Inventor	Goldstein, Joel			
		Examiner Name	Riddick, Marie L.			
Applicant claims small entity status.	See 37 CFR 1.27	Art Unit	1713			
TAL AMOUNT OF PAYMENT	(\$) 320	Attaca and Dandard No.	00070 LICA			

TOTAL A

Attorney Bocket No.   00070 03A						
METHOD OF PAYMENT (check all that apply)	FEE CALCULATION (continued)					
Check Credit card Money Order None	3. ADDITIONAL FEES					
Deposit Account:	I —		Small		1	
Denneit	Fee Code	Fee (\$)	Fee Code	Fee (\$)	Fee Description	Fee Paid
Account 01-0493	1051	130	2051		Surcharge - late filing fee or oath	ree Faiu
Number Deposit Account Air Products and Chemicals, Inc.		50	2052	25	Surcharge - late provisional filing fee or cover sheet	
Name The Director is authorized to: (check all that apply)		130	1053	130	Non-English specification	
Charge fee(s) indicated below Credit any overpayments	1812	2,520	1812	2,520	For filing a request for ex parte reexamination	<u> </u>
Charge any additional fee(s) or any underpayment of fee(s)		920*	1804	920*	Requesting publication of SIR prior to	
Charge fee(s) indicated below, except for the filing fee		4 040*	4005	4 0 40*	Examiner action	
to the above-identified deposit account.		1,840*	1805	1,840	Requesting publication of SIR after Examiner action	
FEE CALCULATION	1251	110	2251	55	Extension for reply within first month	
1. BASIC FILING FEE	1252	420	2252	210	Extension for reply within second month	
Large Entity Small Entity	1253	950	2253	475	Extension for reply within third month	<b></b>
Fee Fee Fee Fee Pee Paid Code (\$) Code (\$)	1254	1,480	2254	740	Extension for reply within fourth month	
1001 770 2001 385 Utility filing fee	1255	2,010	2255	1,005	Extension for reply within fifth month	
1002 340 2002 170 Design filing fee	140.1	330	2401	165	Notice of Appeal	
1003 530 2003 265 . Plant filing fee	1402	330	2402	165	Filing a brief in support of an appeal	320
1004 770 2004 385 Reissue filing fee	1403	290	2403	145	Request for oral hearing	
1005 160 2005 80 Provisional filing fee	1451	1,510	1451	1,510	Petition to institute a public use proceeding	
SUBTOTAL (1) (\$) 0	1452	110	2452	55	Petition to revive - unavoidable	
		1,330	2453	665	Petition to revive - unintentional	
2. EXTRA CLAIM FEES FOR UTILITY AND REISSUE		1,330	2501	665	Utility issue fee (or reissue)	
Extra Claims below Fee Paid		480	2502	240	Design issue fee	
Total Claims	1503	640	2503	320	Plant issue fee	
Claims - 3** = X O	1460	130	1460	130	Petitions to the Commissioner	
	1807	50	1807	7 50	Processing fee under 37 CFR 1.17(q)	
Large Entity   Small Entity Fee Fee Fee Fee Fee Description	1806	180	1806		Submission of Information Disclosure Stmt	
Code (\$) Code (\$)	8021	. 40	802 <sup>-</sup>	1 40	Recording each patent assignment per property (times number of properties)	
1202 18 2202 9 Claims in excess of 20 1201 86 2201 43 Independent claims in excess of 3	1809	770	2809	9 385	Filing a submission after final rejection (37 CFR 1.129(a))	
1203 290 2203 145 Multiple dependent claim, if not paid	1810	770	2810	385	For each additional invention to be	
1204 86 2204 43 ** Reissue independent claims over original patent	1801	770	2801	385	examined (37 CFR 1.129(b))  Request for Continued Examination (RCE)	
1205 18 2205 9 ** Reissue claims in excess of 20 and over original patent	1802	900	1802	900		
SUBTOTAL (2) (\$) 0	Other fee (specify)					
SUBTOTAL (2) (\$) 0 **or number previously paid, if greater; For Reissues, see above	*Redu	ced by	Basic I	Filing F	ee Paid SUBTOTAL (3) (\$)	320

SUBMITTED BY				(Complete	(if applicable))
Name (Print/Type)	Mary E. Bongiomo	Registration No. (Attorney/Agent)	36091	Telephone	610-481-8820
Signature	Mary & Bonga	orno		Date	04/05/04

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DOCKET NO.: 06076 USA

#### IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE APPLI-

CATION OF

Goldstein, et al.

: CONFIRMATION NO.: 6932

**SERIAL NO.** 

09/699,723

: GRP. ART UNIT:

1713

**FILED** 

October 30, 2000

: EXAMINER: Reddick, Marie L.

FOR

REDUCED FORMALDEHYDE NONWOVEN BINDERS WHICH

CONTAIN POLYMERIZED UNITS OF N-METHYLOLACRYLAMIDE

**CUSTOMER NO.:** 

23543

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Sir:

# APPEAL BRIEF UNDER 37 CFR 1.192(b)

This appeal is from the final rejection mailed on May 21, 2003.

### **REAL PARTY IN INTEREST**

Air Products Polymers, L.P. is the real party in interest in the appeal. The assignment has been recorded at Reel/Frame 012762/0076.

## **RELATED APPEALS AND INTERFERENCES**

There are no related appeals or interferences.

#### STATUS OF CLAIMS

Claims 1-7 are pending and are being appealed. Claims 1-7 were rejected in a final office action dated May 21, 2003, and an advisory action mailed on February 3, 2004.

### STATUS OF THE AMENDMENTS

All amendements have been entered.

#### SUMMARY OF INVENTION

The invention relates to use of a special class of reducing agents that is part of a redox catalyst system for the emulsion polymerization of vinyl acetate, N-methylolacrylamide (NMA), and, optionally, another monomer. The reducing agents have the formula

$$\begin{array}{c|c}
O & R_1 \\
S & R_2 \\
R_3 & R_3
\end{array}$$

wherein M is a hydrogen atom, an ammonium atom or a monovalent metal ion,  $R_1$  is OH or  $NR_4R_5$  wherein  $R_4$  and  $R_5$  each are H or  $C_1$ - $C_6$  alkyl;  $R_2$  is H or an alkyl, alkenyl, cycloalkyl or aryl and  $R_3$  is  $CO_2M$ . The preferred reducing agent is the glycolic acid adduct of sodium sulfite which is sold under the trademark Bruggolite FF-6. (page 6, lines 16-18, of the specification) One of the unexpected benefits of using these reducing agents as part of the redox catalyst system is that the polymer emulsion has reduced free formaldehyde content. (page 3, lines 1-14; and page 5, line 22 to page 6, line 18, of the specification)

#### **ISSUES**

There is one rejection presented in the final office action. Claims 1-7 are rejected under 35 U.S.C. 103(a) as being unpatentable over Berghofer et al. (US 6,211,400) in combination with Applicants' disclosure.

#### **GROUPING OF CLAIMS**

Claims 1 through 7 are to be considered as one group.

#### **ARGUMENT**

The Examiner maintained the rejection of Claims 1-7 under 35 USC 103(a) as being unpatentable over Berghofer et al (US 5,211,400 B1) in combination with Applicants' own disclosure. The Examiner stated at page 3 first full sentence of the final rejection (paper number 15):

"It is tenable that the sulfinic acid derivatives (reducing agents) used in the emulsion polymerization technique of Berghofer et al would generate a final polymer emulsion governed by a reduction in formaldehyde since the vinyl acetate based polymer emulsion of Berghofer et al, as modified, is essentially the same as the claimed vinyl acetate based polymer emulsion and there is nothing iron clad on this record diffusing this issue."

In the office action mailed on May 21, 2003 (paper number 13), at page 4, first paragraph, the Examiner referred to the Background of the Invention at page 1, lines 20-21 and page 2, lines 1-15 of Applicants' disclosure as support for modifying the polymer disclosed in Run 11 of Berghofer et al. to include NMA as a co-monomer. However, Berghofer et al. does not disclose or suggest using a crosslinking monomer in the polymers disclosed therein and specifically does not disclose or suggest using NMA as a crosslinking monomer. Use of a crosslinking monomer, such as NMA, produces a distinctly different polymer than the polymers disclosed by Berghofer et al.

It should also be noted that page 1, lines 12-14, of the instant specification, reads: "Reduction of formaldehyde in vinyl acetate based emulsions has been achieved by using less favored reducing agents to the formaldehyde sulfoxylates, or by reducing the level of N-methylol acrylamide. ..."

Since Berghofer et al. teach that the sulfinic acid derivatives disclosed therein have a reducing action similar to formaldehyde sulfoxylates but do not eliminate formaldehyde before, during, and after use, it would not be expected that use of the sulfinic acid derivatives in producing polymers from vinyl acetate, NMA, and optionally another monomer, would result in a reduction in residual formaldehyde.

As the Examiner indicated, the reference is evaluated, as a whole, for what it fairly teaches. It teaches, as a whole, that the novel compounds disclosed therein can be used as reducing agents and will not eliminate formaldehyde before, during and after use. Berghofer et al. do not disclose or suggest polymerization in which residual formaldehyde is formed, such as when NMA is used as a co-monomer.

The ability to use NMA in the polymer and reduce the amount of formaldehyde that would typically be present in the resulting emulsion polymer is the value of the reducing agents of this invention. The formaldehyde-free reducing agent of this invention

unexpectedly reduces the amount of formaldehyde in the final polymer emulsion product derived from vinyl acetate vinyl acetate, NMA and optionally other monomers, compared to other formaldehyde-free reducing agents. See Example 2 of this specification in which polymers of vinyl acetate, ethylene, and NMA were produced using either ascorbic acid (sodium erythorbate) or Bruggolite FF 6 as reducing agent. Use of sodium erythorbate resulted in no additional formaldehyde being formed during the polymerization process. However, unexpectedly, use of Bruggolite FF 6, in the same process, resulted in a substantial reduction of formaldehyde. A summary of the 3 different runs reported in Example 2 is presented below:

	Amount of Formaldehyde (ppm) using One of the following Reducing Agents:			
Vinyl Acetate-Ethylene-NMA Polymer	Sodium Erythorbate	Bruggolite FF 6		
A (Tg = -14 °C)	27.2	3.3		
B (Tg = 10 °C)	57.1	8.6		
C (Tg = 0 °C)	47.5	6.8		

The above described reduction in formaldehyde is not reported by Berghofer et al. In fact, Berghofer et al state, at col. 4, lines 22-24, that the reducing compounds disclosed therein have a reducing action comparable to formaldehyde sulfoxylate; but they do not eliminate formaldehyde before, during, or after use.

The above data rebut a prima facie obviousness rejection based on Berghofer et al together with what is well known and admitted in the Background of this Invention.

It is therefore submitted that the claimed invention would not have been obvious based on Berghofer et al together with what is well known and admitted in the Background of this Invention. Berghofer et al do not teach preparation of polymers containing NMA, and do not teach or suggest that formaldehyde would be lowered during the preparation of polymer emulsions in which reducing agents described therein are used. In addition, the data in this case rebut a prima facie obviousness rejection based on Berghofer et al together with what is well known and admitted in the Background of this Invention.

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Reversal of the rejection is requested.

Respectfully submitted,

Mary & Bongwino
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#### **APPENDIX**

### Claims Involved in the Appeal

1. In a vinyl acetate based polymer emulsion formed by the emulsion polymerization of vinyl acetate and N-methylolacrylamide, optionally other monomers, in the presence of a stabilizing system and a redox catalyst system comprised of an oxidizing agent and a reducing agent, the improvement for reducing formaldehyde emissions in the resulting vinyl acetate based polymer emulsion, which comprises:

forming said vinyl acetate based polymer emulsion utilizing as the reducing component of the redox catalyst system a reducing agent of the formula:

$$\begin{array}{c|c}
O & R_1 \\
S & R_2 \\
R_3 & R_2
\end{array}$$

where M is a hydrogen atom, an ammonium atom or a monovalent metal ion,  $R_1$  is OH or  $NR_4R_5$  wherein  $R_4$  and  $R_5$  each are H or  $C_1$ - $C_6$  alkyl;  $R_2$  is H or an alkyl, alkenyl, cycloalkyl or aryl and  $R_3$  is  $CO_2M$ .

- 2. The vinyl acetate based polymer emulsion of Claim 1 in which the vinyl acetate based polymer comprises polymerized units of ethylene in an amount of from about 10 to 40% by weight of the polymer.
- 3. The vinyl acetate based polymer emulsion of Claim 2 wherein the N-methylolacrylamide is present in an amount of from about 0.5 to 10% by weight of the polymer.
- 4. The vinyl acetate based polymer emulsion of Claim 3 wherein the reducing agent represented by the formula is selected from the group consisting of: 2-hydroxyphenyl hydroxymethyl sulfinic acid-sodium salt; 4-methoxyphenyl hydroxymethyl sulfinic acid-sodium salt; 2-hydroxy-2-sulfinato acetic acid-disodium salt; 2-hydroxy-2-sulfinato acetic acid-zinc

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salt; 2-hydroxy-2-sulfinato propionate-disodium salt; ethyl 2-hydroxy-2-sulfinato propionate-sodium salt.

- 5. The vinyl acetate based polymer emulsion of Claim 3 wherein the vinyl acetate based polymer emulsion is formed using a redox catalyst system of hydrophobic hydroperoxide and the glycolic acid adduct of sodium sulfite.
  - 6. The vinyl acetate based polymer emulsion of Claim 3 wherein M is sodium or zinc.
  - 7. The vinyl acetate based polymer emulsion of Claim 3 wherein  $R_1$  is OH.